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or aspartate or salicylate)
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         53909 ASPARTATE
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               OR PYRUVATE OR SUCCINATE OR ADIPATE OR ASPARTATE OR SALICYLATE)
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## => d bib abs kwic ANSWER 1 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN 2002:521462 CAPLUS AN DN137:88442 Incensole and furanogermacrens and compounds in treatment for inhibiting neoplastic lesions and microorganisms Shanahan-Pendergast, Elisabeth ΤN PΑ Ire. SO PCT Int. Appl., 68 pp. CODEN: PIXXD2 рΤ Patent English LA FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE \_\_\_\_\_\_ \_\_\_\_\_ WO 2002053138 A2 20020711 WO 2002-IE1 20020102 PΙ WO 2002053138 20020919 **A**3 W: AE, AG, AT, AU, BB, BG, CA, CH, CN, CO, CU, CZ, LU, LV, MA, MD, UA, UG, US, VN, YU, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, AT, BE, CH, CY, DE, ES, FI, ML, MR, NE, SN, TD, TG A2 20031015 EP 2002-727007 20020102 EP 1351678 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR US 2004092583 A1 20040513 US 2004-250535 20040102 PRAI IE 2001-2 20010102 Α 20020102 WO 2002-IE1 MARPAT 137:88442 The invention discloses the use of incensole and/or furanogermacrens, derivs. metabolites and precursors thereof in the treatment of neoplasia, particularly resistant neoplasia and immundysregulatory disorders. These compds. can be administered alone or in combination with conventional chemotherapeutic, antiviral, antiparasite agents, radiation and/or surgery. Incensole and furanogermacren and their mixture showed antitumor activity against various human carcinomas and melanomas and antimicrobial activity against Staphylococcus aureus and Enterococcus faecalis. 50-07-7, Mutamycin 50-18-0, Cyclophosphamide 50-28-2, Estradiol, 50-35-1, Thalidomide 50-76-0, Dactinomycin biological studies 50-91-9, Floxuridine 51-21-8, Fluorouracil 51-75-2, Mechlorethamine 52-24-4, Thiotepa 53-19-0, Mitotane 53-43-0, DHEA 53-79-2, Puromycin 54-71-7, Pilocarpine hydrochloride 54-91-1, Pipobroman 55-21-0D, Benzamide, N-substituted compds. 55-86-7, Mechlorethamine Hydrochloride 55-86-7D, Nitrogen mustard, derivs. 55-98-1, Busulfan 56-53-1, Diethylstilbestrol 57-22-7, Vincristine 57-63-6, Ethinyl oestradiol 57-83-0, Progesterone, biological studies 58-05-9, Leucovorin 58-58-2, Puromycin Hydrochloride 59-05-2, Methotrexate 66-75-1, Uracil Mustard 83-89-6, Acriquine 101-60-0, Porphyrin 106-60-5, Aminolevulinic acid 114-70-5, Sodium phenylacetate 122-79-2, Phenylacetate 125-45-1, Azetepa 125-84-8, Aminoglutethimide 127-07-1, Hydroxyurea 143-67-9, Vinblastine Sulfate 145-63-1, Suramin 147-94-4, Cytarabine 148-82-3, Melphalan 154-42-7, Thioguanine 154-93-8, Carmustine 302-49-8, Uredepa 302-79-4, Tretinoin 305-03-3, Chlorambucil 320-67-2, Azacitidine 359-83-1, Pentazocine 364-62-5, Metoclopramide 366-70-1, Procarbazine Hydrochloride 378-44-9, Betamethasone 423-55-2, Perflubron 459-86-9, Mitoguazone 465-65-6, Naloxone 472-15-1, Betulinic acid 481-29-8, Epiandrosterone 518-28-5, Podophyllotoxin 520-85-4, Medroxyprogesterone 521-12-0, Dromostanolone Propionate 536-59-4, Perillyl alcohol 548-04-9, Hypericin 566-48-3, Formestane 578-95-0D, Acridone, imidazo derivs. 578-95-0D, Acridone, propylbis derivs. 595-33-5, Megestrol Acetate 645-05-6, Altretamine 646-08-2, β-Alethine 671-16-9, Procarbazine 801-52-5, Porfiromycin 865-21-4, Vinblastine 911-45-5, Clomifene 968-93-4, Testolactone 1271-19-8, Titanocene dichloride 1402-81-9, Ambomycin 1403-99-2, Mitogillin 1404-00-8, Mitomycin 1404-15-5. Nogalamycin 1404-20-2, Peliomycin 1404-64-4, Sparsomycin 1661-29-6, Meturedepa 1972-08-3, Dronabinol 1980-45-6, Benzodepa 2068-78-2, Vincristine Sulfate 2353-33-5, Decitabine 2508-89-6 2608-24-4, Piposulfan 2809-21-4D, Etidronic acid, rhenium-186 complexes 2919-66-6, Melengestrol acetate 2998-57-4, Estramustine 2998-57-4D, Estramustine, analogs 3073-59-4, Hexamethylene bisacetamide 3094-09-5, Doxifluridine 3562-63-8, Megestrol 3778-73-2, Ifosfamide 3930-19-6, Streptonigrin 4105-38-8 4291-63-8, Cladribine 4342-03-4, Dacarbazine 4342-07-8 4803-27-4, Anthramycin 5072-26-4, Buthionine sulfoximine 5373-42-2, Thaliblastine 5508-58-7, Andrographolide 5579-27-1, Simtrazene 5581-52-2, Thiamiprine 5696-17-3, Epipropidine 6157-87-5,

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Platinum, lipophilic compds. or complexes

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RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
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ANSWER 2 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN
AN
    1992:99301 CAPLUS
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     Maleic anhydride copolymers as antidotes for the cytotoxicity of neoplasm
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     G.D. Searle and Co., USA
PA
SO
    Eur. Pat. Appl., 27 pp.
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$$\begin{array}{c} \text{Me} \\ \text{C}_{6}\text{H}_{5} - \text{CH} & \text{CH} - \text{CH} - \text{CH}_{2} - \text{CH}_{2} \\ \text{CH} - \text{CH} - \text{CH}_{2} - \text{CH}_{2} \\ \text{C}_{-0} & \text{C}_{-0} \\ \text{NH}_{2} & \text{O}_{-1} \\ \text{a} & \text{NH}_{4}^{+} \\ \end{array} \right] \begin{array}{c} \text{CH} - \text{CH}_{2} - \text{CH}_{2} \\ \text{C}_{-0} & \text{C}_{-0} \\ \text{NH}_{2} & \text{O}_{-1} \\ \text{NH}_{2} & \text{O}_{-1} \\ \text{NH}_{4}^{+} \\ \end{array} \right)$$

Half-amide:half-imide copolymers comprising ethylene and maleic anhydride moieties (structure given), specifically carbetimer (I; a/b = 1:2-5), decrease the cytotoxic side effects of neoplasm inhibitors. Mice treated i.v. with 21 mg adriamycin/kg died within 5 days. When 1700 mg I/kg was administered concomitantly, no lethality was shown for >30 days. 50-18-0, Cyclophosphamide 50-76-0, Dactinomycin 50-91-9, Floxuridine 51-21-8, 5-Fluorouracil 51-21-8D, conjugates with fibrinogens 53-19-0, Mitotane 54-42-2, NSC 39661 56-18-8, Norspermidine 57-22-7, Vincristine 59-05-2, Methotrexate 75-19-4D, Cyclopropane, spiro derivs. 113-15-5, ERgotamine 127-07-1 143-67-9, Vinblastine sulfate 147-94-4, Cytarabine 147-94-4D, Cytarabine, conjugates 154-42-7, Thioguanine 154-93-8, Carmustine 302-79-4, Retinoic acid 305-03-3, Chlorambucil 432-70-2, α-Carotene 636-65-7, Isoglutamine 642-18-2, Alstonine 645-05-6, Altretamine 671-16-9, Procarbazine 1149-99-1, Illudin 1404-00-8, Mitomycin 1404-64-4, Sparsomycin 1948-56-7D, Dehydroalanine, N-acyl derivative 2353-33-5, NSC 127716 3073-59-4, NSC 95580 3094-09-5, Doxifluridine 3778-73-2, Ifosfamide 4005-51-0, Aminothiadiazole 4342-03-4 4759-48-2, Isotretinoin 5373-42-2, Thaliblastine 6620-60-6, Proglumide 6829-55-6 7440-06-4D, Platinum, derivs., complexes 7481-89-2, Dideoxycytidine 7534-61-4, NSC 145813 9014-02-2D, Neocarzinostatin, conjugates with styrene-maleci acid copolymer 9015-68-3, Asparaginase 9041-93-4, Bleomycin sulfate 9054-89-1, Superoxide dismutase 10318-26-0, Mitolactol 12633-27-1, T 680 13010-47-4, Lomustine 13494-90-1, Gallium nitrate 13665-88-8, Mopidamol 13909-02-9 13909-09-6, Semustine 14459-29-1D, polymers 14930-96-2, Cytochalasin B 15219-97-3, Oxalysine 15663-27-1, Cisplatin 18378-89-7, Plicamycin 19624-67-0, SKF 101772 20830-81-3 21416-67-1, Razoxane 22862-76-6, Anisomycin 23214-92-8, Doxorubicin 23214-92-8D, conjugates with fibrinogens 24584-09-6, ICRF 187 25300-64-5D, conjugates with neocarzinostatin 26833-87-4, Homoharringtonine 27686-84-6, CHX 100 28656-91-9D, Aeroplysinin, derivs. 29069-24-7, Prednimustine 29767-20-2, Teniposide 31430-18-9D, Nocodazole, N-acyl derivative 33069-62-4, Taxol 33419-42-0, Etoposide 35144-64-0D,

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Aldophosphamide, analogs 38077-12-2 39389-47-4, Distamycin
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Estramustine phosphate sodium 53123-88-9, Rapamycin 53643-48-4,
Vindesine 53910-25-1, Pentostatin 54083-22-6, Zorubicin 54350-48-0,
Etretinate 54526-94-2, Steffimycin B 54824-17-8, Mitonafide
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60784-46-5, Elmustine 61251-97-6, Baccharin 61422-45-5, Carmofur
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69408-81-7, Amonafide 69772-39-0, Neoenactin 69839-83-4, Didox
70189-62-7, TA 077
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95604-83-4, Probimane 95693-76-8, DATHF 95734-82-0, 254S 96086-68-9,
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RL: PRP (Properties)
    (cytotoxicity of, maleic anhydride copolymer antidote for)
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$$\begin{array}{c} R^1 \\ 0 \\ N \\ \end{array} \qquad \qquad \qquad HX$$

AB Salts of amonafide or amonafide analogs I [R1 = (un)substituted NH2, aminoalkyl; R2 = OH, alkoxy, (un)substituted NH2, SO3H, NO2, acyloxy; X = carboxylate] were prepared Thus, 3-nitro-1,8-naphthalic anhydride was treated with Me2NCH2CH2NH2, followed by L-malic acid to give mitonafide malate which was reduced over Pd/C to give amonafide malate. This compound was completely soluble in H2O and normal saline solution and had anticancer activity both in vitro and in vivo.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

AB Salts of amonafide or amonafide analogs I [R1 = (un) substituted NH2, aminoalky1; R2 = OH, alkoxy, (un) substituted NH2, SO3H, NO2, acyloxy; X = carboxylate] were prepared Thus, 3-nitro-1,8-naphthalic anhydride was treated with Me2NCH2CH2NH2, followed by L-malic acid to give mitonafide malate which was reduced over Pd/C to give amonafide malate. This compound was completely soluble in H2O and normal saline solution and had anticancer activity both in vitro and in vivo.